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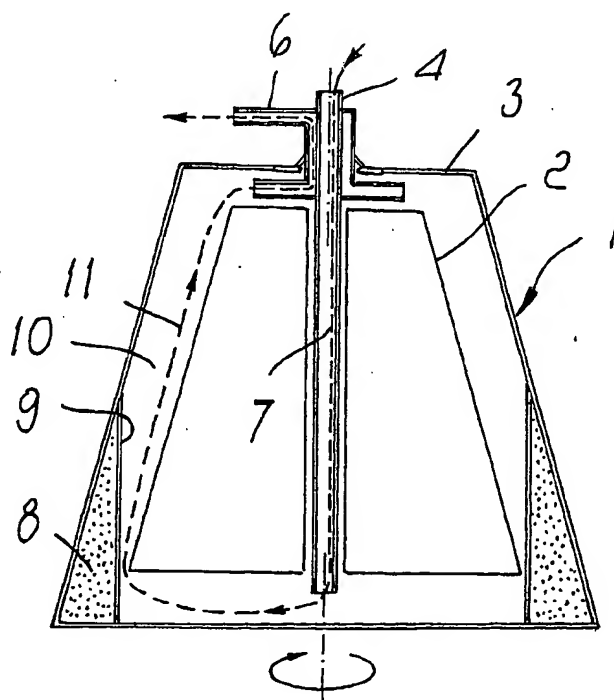
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(54) **Method and apparatus for controlling the washing step in a blood centrifugation cell**

(57) A method and apparatus for controlling the washing step in a blood centrifugation cell in which washing solution is introduced into the blood centrifugation cell and the cell contains compacted red cells and

supernatant at the beginning of the washing step. The apparatus comprises a computer that executes an algorithm that produces a first output that is the concentration of the supernatant in the supernatant-washing solution mixture.

Fig.1



Description

[0001] The invention relates to blood centrifuges and in particular relates to a system for controlling the washing step in a blood centrifugation cell.

[0002] It is known that in some medical procedures, such as inter- and postoperative autotransfusion, there is the need to separate the plasma from the red cells of the blood aspirated from the operating area, so as to make them available for re-infusion to the patient. It is also known that currently this procedure is performed in centrifugation cells in which the blood is introduced by means of a peristaltic pump.

[0003] A centrifugation cell substantially comprises two bells which are mutually coaxial and rigidly coupled, and the portion of space between them is connected to the outside by means of two tubes, an inlet tube and an outlet tube, which are connected to the bells by means of a rotary coupling. The two bells are turned about their own axis, while the tubes are kept motionless.

[0004] The procedure provides for a first step for filling the cell, in which the blood is introduced by means of the inlet tube. Due to the centrifugal force, the red cells, which are the heaviest cellular components of blood, are propelled outward, compacting against the wall of the outer bell. Other cellular components such as white blood cells and platelets are arranged in a thin layer known as buffy coat directly adjacent to the mass of compacted red blood cells. The separated plasma, the remaining component of blood, is arranged in a layer which lies above the buffy coat. The plasma, which contains various substances such as anticoagulant, free hemoglobin and other substances from the operating field, will be referenced hereinafter as "supernatant".

[0005] As filling continues, the buffy coat moves closer to the rotation axis, displacing the supernatant toward the outlet tube of the cell. When the supernatant reaches the outlet tube the supernatant flows out of the cell into an adapted collection bag.

[0006] The flow of the supernatant in the outlet tube continues until an optical detector reports that the buffy coat has reached the outlet tube of the cell. At this point the filling step has ended and the introduction of new blood into the cell ends. The cell now contains compacted red cells and supernatant, which must be eliminated since it cannot be re-infused to the patient together with the red cells.

[0007] The above-described filling step is followed by a washing step performed by means of a washing solution which, when introduced into the cell, gradually takes the place of the supernatant that is expelled. At the end of the washing step the cell contains red cells and washing solution, i.e., substances suitable to be re-infused to the patient. The contents of the cell are collected in a suitable bag in a third step of the procedure, known as emptying.

[0008] Our attention is focused exclusively on the washing step, which is currently performed in manners

that are not entirely satisfactory. A first procedure adopted in the background art provides for introducing in the cell a preset amount of washing solution at a value that is assuredly more than sufficient to wash the supernatant. The consequent oversizing, however, wastes time and washing solution.

[0009] Another procedure used in the art provides a transparency sensor on the outlet duct. However, this sensor is not able to detect the passage of transparent components of the supernatant such as the anticoagulant, and therefore does not provide entirely satisfactory results.

[0010] The invention provides a system for controlling the washing step in which the washing step can be stopped when the intended result is reached. The present invention provides a method for controlling a washing step in a blood centrifugation cell wherein washing solution is introduced into the blood centrifugation cell in the washing step, the cell containing compacted red blood cells and supernatant at the beginning of the washing step. The method comprises: (1) providing a blood centrifugation cell, a pump for communicating liquid to the blood centrifugation cell, and a computer configured to receive data and produce at least one output; (2) providing first input data to the computer indicative of the total volume of liquid that has entered the cell during the filling step that precedes the washing step and during the washing step itself, the liquid being blood during the filling step and the liquid being washing solution during the washing step; (3) providing second input data to the computer indicative of the hematocrit value of the blood that was introduced during the filling step; (4) providing third input data to the computer indicative of the geometric characteristics of the cell; (5) processing the first, second, and third input data in the computer to produce a first output, said first output being the concentration of the supernatant in the supernatant-washing solution mixture that is present in the cell during the washing step, said first output being produced by the computer executing an algorithm that expresses the supernatant extinction law inside the cell using the first, second, and third input data; and (6) stopping the washing step when a certain concentration of supernatant in the supernatant-washing solution is reached.

[0011] In addition, the invention provides an apparatus for performing the method for controlling the washing step in a blood centrifugation cell.

Figure 1 is a schematic view of a blood centrifugation cell.

Figure 2 is a diagram of an apparatus of the invention.

[0012] With reference to the figures, the reference numeral 1 generally designates the blood centrifugation cell, which comprises inner bell 2 and outer bell 3, which are mutually rigidly coupled and are made to rotate in the direction of the arrow in the figures. The numeral 4

respectively designates the inlet tube to the portion of space comprised between the two bells. Peristaltic pump 5 pumps fluids into inlet tube 4. The fluids follow path 7 into region 10 between the inner and outer bells. Centrifugation separates blood components, as described further below, and supernatant flows out of the centrifuge via outlet tube 6. The inlet and outlet tubes are connected to the assembly of the inner and outer bells by means of a rotary coupling, so that they can remain motionless.

[0013] During the cell filling step, the red blood cells enter the cell along path 7 due to the action of the peristaltic pump 5, which is connected at the suction to a container known as a cardiotomy reservoir, not shown in the figures. During centrifugation, the red blood cells are compacted in region 8, and the supernatant follows path 11 to the outlet tube 6. Buffy coat 9 separates the compacted red blood cells from the supernatant in region 10. The supernatant then flows toward the outlet tube 6 of the cell along the path 11.

[0014] When buffy coat 9, by moving increasingly closer to the rotation axis, reaches the full level indicated by a sensor, the introduction of blood into the cell ceases as pump 5 stops. Now the filling step has ended and cell 1 contains compacted red cells and supernatant.

[0015] This is followed by a washing step to eliminate the supernatant by means of a washing solution. Preferred washing solution consists of a physiological saline solution 0.9g/l NaCl/water for generic medical use. The washing solution is conveyed to cell 1 through inlet tube 4 by pump 5, which is in communication with a reservoir of washing solution.

[0016] The washing solution gradually takes the place of the supernatant, and at the end of the washing step in cell 1, the replacement of the supernatant with the washing solution has occurred substantially completely. A minute amount of supernatant in cell 1 remains and will be reinfused to the patient. However, small amounts of supernatant are obviously tolerable.

[0017] During the washing step, therefore, in the volume of the cell 1 that is not occupied by the compacted red cells, there is a mixture of supernatant and washing solution. The expression "supernatant concentration" is used to designate the ratio between the volume of supernatant present in said mixture and the total volume of said mixture, and it is immediately evident that the value of said concentration varies during washing from the initial value of 1, when all the space available is occupied by the supernatant, toward the ideal final value, which is zero and would be reached if the supernatant were eliminated completely and fully replaced by the washing solution. The expression "supernatant extinction law" is used to designate the law that regulates the variation of the concentration of supernatant in the supernatant-washing solution mixture as it decreases from the initial value of 1 toward the final value.

[0018] The control system includes sensor 12, which is suitable to provide the hematocrit reading of the blood

entering the cell during the filling step, encoder or sensor 13 on the driving shaft of the peristaltic pump 5, which detects data related to the rotation angles of said shaft, and computer 14.

[0019] The computer executes an algorithm, derived from a mathematical model or from the processing of experimental data, that expresses the supernatant extinction law within cell 1 and has three inputs and one output.

[0020] The first input comprises the volume of the liquid that enters the cell during the filling step, which is blood, and during the washing step, which is washing solution. This first input is provided, in the described embodiment, by the encoder 13. The data it transmits to the computer 14 related to the rotation angles gradually covered by the pump 5 are converted, since the characteristics of said pump and of the tube 4 are known, into data related to the volume of liquid progressively conveyed. However, clearly the encoder 13 might be replaced with any liquid flow measurement instrument.

[0021] The second input comprises the hematocrit reading of the blood entering cell 1 during the filling step, and is provided by sensor 12, which reports to computer 14 the hematocrit reading of the individual small volumes of blood that enter cell 1 continuously. The second input might be provided in other forms. For example, the computer 14 could include an operator interface which allows entering into the computer 14 data related to the hematocrit value of the blood, which can be determined from the cardiotomy reservoir located at the intake of the pump 5.

[0022] The third input comprises the geometric characteristics of the cell. Accordingly, there are means that allow the operator to enter into computer 14 data related to these characteristics. Alternatively, a sensor could be provided for automatic detection of said characteristics.

[0023] On the basis of the three listed inputs, computer 14 provides an output at each instant of the value of the concentration of supernatant in the supernatant-washing solution mixture that is present in the cell 1 during the washing step. In addition, it is possible to provide time as a fourth input.

[0024] The invention includes means that allow stopping the washing step when the intended conditions are reached. In the described embodiment there is the display 14a, which shows at each instant the value of the concentration of supernatant and thus allows the operator to intervene and turn off the pump 5 when said value reaches the threshold deemed acceptable. According to a different embodiment, there is a controller to stop automatically the operation of the pump 5 when said concentration reaches the threshold value that is preset as acceptable.

[0025] Various modifications and alterations to this invention will become apparent to those skilled in the art without departing from the scope and spirit of this invention. It should be understood that this invention is not intended to be unduly limited by the illustrative embod-

lements and examples set forth herein and that such examples and embodiments are presented by way of example only with the scope of the invention intended to be limited only by the claims set forth herein as follows.

[0026] The disclosures in Italian Patent Application No. MI2001A000899 from which this application claims priority are incorporated herein by reference.

[0027] Where technical features mentioned in any claim are followed by reference signs, those reference signs have been included for the sole purpose of increasing the intelligibility of the claims and accordingly, such reference signs do not have any limiting effect on the interpretation of each element identified by way of example by such reference signs.

Claims

1. A method for controlling a washing step in a blood centrifugation cell wherein washing solution is introduced into the blood centrifugation cell in the washing step, the cell containing compacted red blood cells and supernatant at the beginning of the washing step, the method comprising:
 - providing a blood centrifugation cell, a pump for communicating liquid to the blood centrifugation cell, and a computer configured to receive data and produce at least one output;
 - providing first input data to the computer indicative of the total volume of liquid that has entered the cell during the filling step that precedes the washing step and during the washing step itself, the liquid being blood during the filling step and the liquid being washing solution during the washing step;
 - providing second input data to the computer indicative of the hematocrit value of the blood that was introduced during the filling step;
 - providing third input data to the computer indicative of the geometric characteristics of the cell;
 - processing the first, second, and third input data in the computer to produce a first output, said first output being the concentration of the supernatant in the supernatant-washing solution mixture that is present in the cell during the washing step, said first output being produced by the computer executing an algorithm that expresses the supernatant extinction law inside the cell using the first, second, and third input data; and
 - stopping the washing step when a certain concentration of supernatant in the supernatant-washing solution is reached.
2. The method of claim 1, wherein the algorithm has time as a fourth input data.
3. The method of one of claims 1 to 2, wherein the algorithm that expresses the supernatant extinction law inside the cell is derived from a mathematical model.
4. The method of one of claims 1 to 2, wherein the algorithm that expresses the supernatant extinction law inside the cell is derived from an experimental analysis.
5. The method of one of claims 1 to 4, wherein the pump is a peristaltic pump comprising on the drive shaft of the pump a sensor suitable to transmit to the computer said first input data related to the rotation angles of said drive shaft.
6. The method of one of claims 1 to 5, further comprising providing a sensor for measuring the volume of liquid entering the cell and suitable to transmit to the computer said first input data.
7. The method of one of claims 1 to 6, further comprising providing a sensor for measuring the hematocrit value of the blood entering the cell during the filling step and suitable to transmit to the computer said second input data.
8. The method of one of claims 1 to 7, further comprising providing an operator interface for entering into the computer said second input data indicative of the hematocrit value of the blood.
9. The method of claim 8, wherein said pump comprises an intake and a cardiotomy reservoir is located at the intake for said pump, and wherein said data indicative of the hematocrit value of the blood is determined by measuring a hematocrit value of the blood in the cardiotomy reservoir.
10. The method of one of claims 1 to 9, further comprising providing an operator interface for entering into the computer said third input data indicative of the geometric characteristics of the cell.
11. The method of one of claims 1 to 10, further comprising providing a sensor for automatically detecting the geometric characteristics of the cell and suitable to transmit to the computer said third input data.
12. The method of one of claims 1 to 11, further comprising providing an operator interface for displaying the concentration of the supernatant in the supernatant-washing solution mixture.
13. The method of claim 12, wherein the operator stops the washing step when a certain concentration of supernatant is reached.

14. The method of one of claims 1 to 12, further comprising providing a controller for stopping the washing step when a certain concentration of supernatant in the supernatant-washing solution is reached.
15. The method of claim 14, wherein the controller stops the washing step when a preset value of the concentration of supernatant in the supernatant-washing solution mixture is reached.
16. An apparatus for controlling a washing step in a blood centrifugation cell wherein washing solution is introduced into the blood centrifugation cell in the washing step, the cell containing compacted red blood cells and supernatant at the beginning of the washing step, the apparatus comprising:
- a pump for communicating liquid to the blood centrifugation cell;
 - a first sensor configured to produce first input data indicative of the total volume of liquid that has entered the cell during the filling step that precedes the washing step and during the washing step itself, the liquid being blood during the filling step and the liquid being washing solution during the washing step;
 - a computer which executes an algorithm that expresses the extinction law of the supernatant inside the cell using said first input data, second input data indicative of the hematocrit value of the blood that was introduced during the filling step, and third input data indicative of the geometric characteristics of the cell, to produce a first output, said first output being the concentration of the supernatant in the supernatant-washing solution mixture;
 - an operator interface for entering input data and displaying said first output data; and
 - a controller for stopping the washing step when a certain concentration of supernatant in the supernatant-washing solution is reached.
17. The apparatus of claim 16, further comprising a second sensor for measuring the hematocrit value of the blood entering the cell during the filling step and suitable to transmit to the computer said second input data.
18. The apparatus of one of claims 16 or 17, further comprising a second sensor for automatically detecting the geometric characteristics of the cell and suitable to transmit to the computer said third input data.
19. The apparatus of one of claims 16 to 18, wherein the pump is a peristaltic pump comprising on the drive shaft of the pump the first sensor suitable to transmit to the computer said first input data related to the rotation angles of said drive shaft.
20. The apparatus of one of claims 16 to 19, further comprising a second sensor for measuring the volume of liquid entering the cell and suitable to transmit to the computer said first input data.

Fig.2

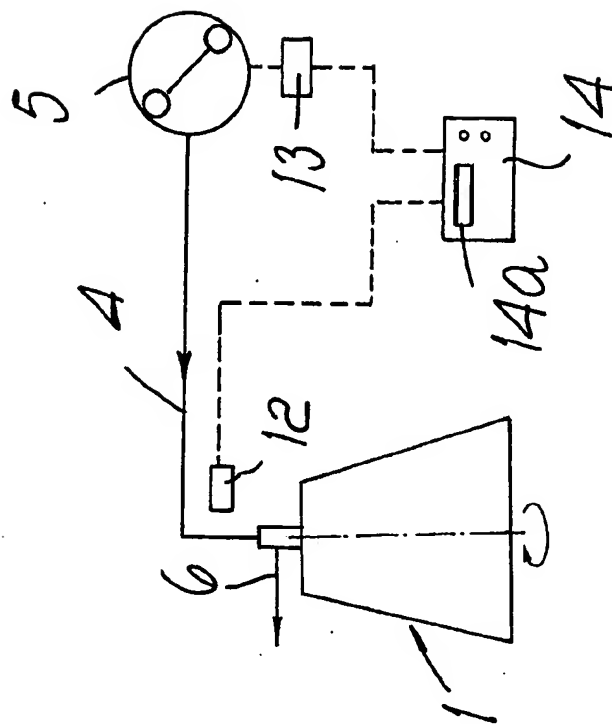
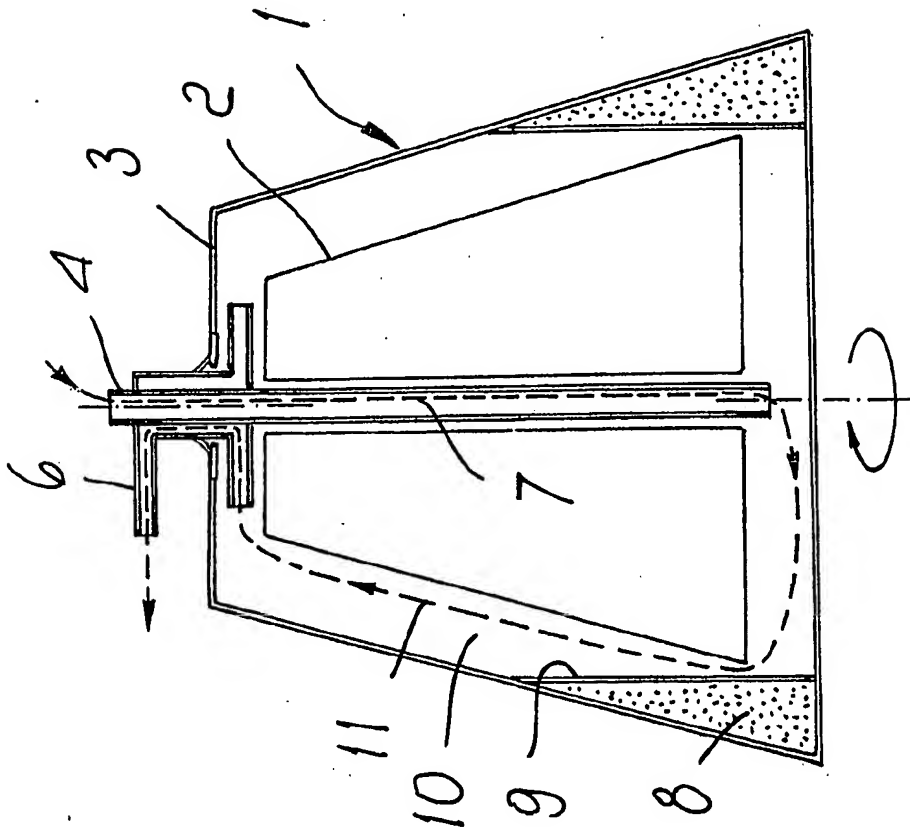


Fig.1





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EUROPEAN SEARCH REPORT

Application Number
EP 02 00 9205

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
A	EP 0 682 953 A (HAEMONETICS CORP) 22 November 1995 (1995-11-22) * page 2, line 53 - page 3, line 2 * * page 3, line 20 - line 57 * ---	1,16	A61M1/36 B04B5/04
A	EP 0 931 554 A (FRESENIUS AG) 28 July 1999 (1999-07-28) * page 2, line 47 - page 3, line 32 * ---		
A	WO 98 29149 A (SHETTIGAR U RAMAKRISHNA) 9 July 1998 (1998-07-09) * page 4, line 1 - page 5, line 26 * -----		
			TECHNICAL FIELDS SEARCHED (Int.Cl.7)
			A61M B04B
The present search report has been drawn up for all claims			
Place of search MUNICH		Date of completion of the search 22 July 2002	Examiner Péru, L
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**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 02 00 9205

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22-07-2002

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 0682953	A	22-11-1995	US 5478479 A	26-12-1995
			EP 0682953 A1	22-11-1995
			JP 3215597 B2	09-10-2001
			JP 7313587 A	05-12-1995
EP 0931554	A	28-07-1999	DE 19802321 A1	05-08-1999
			EP 0931554 A2	28-07-1999
			JP 11267195 A	05-10-1999
			US 6299784 B1	09-10-2001
			US 2002011453 A1	31-01-2002
WO 9829149	A	09-07-1998	US 5876611 A	02-03-1999
			WO 9829149 A1	09-07-1998

EPO FORM P445

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